Abstract Accelerating progress towards the United Nations' 90-90-90 target: #89 The impact of a province-wide HIV Treatment-as-Prevention-based initiative in British Columbia, Canada

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Introduction

"HIV Treatment as Prevention" (TasP), the scaling-up of testing followed by the immediate initiation of ART, is a strategy for reducing AIDS-related morbidity and mortality, and the spread of HIV. In British Columbia (BC), Canada, TasP was implemented under the Seek and Treat for Optimal Prevention of HIV/AIDS initiative (STOP) starting in 2010.

Objective: To compare the time from HIV diagnosis to antiretroviral therapy (ART) initiation (time Dx-Tx), and from ART initiation to first virologic suppression (time Tx-Vx) before and after the implementation of STOP.



Methods

Design: population-based retrospective cohort study

Data: longitudinal individual-level from STOP cohort

Study population: all diagnosed people living with HIV (PLWH) in BC, who were ≥18 years old, ART naïve, and newly diagnosed in BC between 2005 and 2016

Outcomes: time Dx-Tx & time Tx-Vx

Exposures: HIV diagnosis & ART initiation eras, each grouped into pre-STOP (2005-2009) and post-STOP (2010-2016)

Statistical analysis: negative binomial regressions modelled the effect of STOP on the time Dx-Tx and time Tx-Vx, adjusting for confounders

Figure 1. The distribution of time Dx-Tx and time Tx-Vx (in months) among PLWH in BC from 2005-2016

Results

- PLWH diagnosed before (N=1601) and after STOP HIV/AIDS (N=1700) were significantly different, e.g., 30% vs. 15% ever injected drugs, and the median CD4 level at diagnosis 280 vs. 380 cells/ μ L
- From 2005 to 2016, median time Dx-Tx and time Tx-Vx **decreased substantially** (*Figure 1*)
- After STOP, time Dx-Tx decreased by as much as 22 months among PLWH aged <30 years, but time Tx-Vx remained the longest among PLWH who live in BC's most rural health authority and had history of injection drug use (Figure 2)

Controlling for confounders including changes in ART eligibility and first-line ART preferences, STOP was associated with a 65% shorter time Dx-Tx (adjusted mean ratio: 0.35 [95%CI: 0.32-0.38]) and a **22% shorter time Tx-Vx** (0.78 [0.72-0.85])

Discussion

In our large population-based cohort with universal health TasP-based coverage, а intervention was significantly associated with early ART initiation and faster time to virologic suppression, thus accelerating progress towards the United Nations' **90-90-90 target**.

Our results support the **global** and equitable expansion of TasP to accelerate the control HIV/AIDS, as currently of recommended by the United Nations.





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Figure 2. The distribution of time from HIV diagnosis to ART initiation and from ART initiation to viral suppression (in months) before and after STOP HIV/AIDS roll-out, stratified by selected demographic and clinical characteristics

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